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100. The composition of claim ⁹⁷97, wherein said anti-diabetic agent comprises a type of anti-diabetic agent selected from the group consisting of thiazolidinediones, sulfonylureas, benzoic acid derivatives, and alpha-glucosidase inhibitors.

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101. The composition of claim ⁹⁹100, wherein said anti-diabetic agent is metformin.

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102. The composition of claim ¹⁰⁰101, wherein the amount of metformin is in the range of about 100 mg up to about 2550 mg per dose.

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103. The composition of claim ⁹⁹100, wherein said anti-diabetic agent is a sulfonylurea.

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104. The composition according to claim ¹⁰²103, wherein said sulfonylurea is acetohexamide, chlorpropamide, tolazimide, tolbutamide, glycazide, glipizide, glyburide, or glimeperide.

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105. The composition according to claim ⁹⁹100, wherein said anti-diabetic agent is a thiazolidinedione.

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106. The composition according to claim ¹⁰⁴105, wherein said thiazolidinedione is troglitazone, rosiglitazone, or pioglitazone.

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107. The composition according to claim ⁹⁹100, wherein said anti-diabetic agent is an alpha-glucosidase inhibitor.

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108. The composition according to claim ¹⁰⁶107, wherein said alpha-glucosidase inhibitor is acarbose or miglitol.

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109. The composition according to claim ¹⁰⁷108, wherein said anti-diabetic agent is a benzoic acid derivative.

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110. The composition according to claim ¹⁰⁸109, wherein said benzoic acid derivative is repaglinide.

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111. The composition of claim ⁹⁷97, wherein said bioavailable source of chromium comprises more than 300 micrograms elemental chromium.

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112. The composition of claim 97, wherein said bioavailable source of vanadium is vanadyl sulfate.

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113. The composition of claim 97, wherein said bioavailable source of vanadium comprises more than about 10 mg elemental vanadium.

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114. The composition of claim 97, further comprising an effective amount of a bioavailable source of one or more of the following: magnesium, and aspirin.

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115. The composition of claim ¹¹¹112, wherein said bioavailable source of chromium is chromium polynicotinate.

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116. A method for improving glucose metabolism, comprising treating a patient for at least about a thirty day period by administering a pharmaceutical composition comprising an anti-diabetic

agent other than insulin, a bioavailable source of chromium, and a bioavailable source of vanadium, wherein the Hb1Ac level for said patient is reduced by at least about 10% after such treatment as compared to treatment with said anti-diabetic agent alone.

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~~117~~. The method of claim ¹¹⁵~~116~~, wherein said anti-diabetic agent comprises a type of anti-diabetic agent selected from the group consisting of thiazolidinediones, sulfonylureas, benzoic acid derivatives, and alpha-glucosidase inhibitors.

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~~118~~. The method of claim ¹¹⁶~~117~~, wherein said anti-diabetic agent is metformin.

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~~119~~. The method of claim ¹¹⁵~~116~~, wherein said bioavailable source of chromium comprises more than 300 micrograms elemental chromium when said composition is administered on a daily basis.

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~~120~~. The method of claim ¹¹⁸~~119~~, wherein said bioavailable source of vanadium comprises at least about 10 mg elemental vanadium when said composition is administered on a daily basis.

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~~121~~. The method of claim ¹¹⁸~~119~~, wherein said bioavailable source of vanadium is vanadyl sulfate.

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~~122~~. The method of claim ¹²⁰~~121~~, further comprising an effective amount of a bioavailable source of one or more of the following: magnesium, and aspirin.

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~~123~~. An ingestible formulation for improving glucose metabolism in a subject with abnormal glucose metabolism, comprising:

(a) a bioavailable source of chromium in a complex and amount that delivers an effective amount of chromium for improving glucose metabolism;

(b) a bioavailable source of vanadium in a complex and amount that delivers an effective amount of vanadium for improving glucose metabolism; and

(c) an anti-diabetic agent other than insulin,

wherein treatment with said composition for at least about thirty days reduces the Hb1Ac level for said subject by at least about 50% as compared to treatment with said anti-diabetic agent alone.

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~~124~~. The ingestible formulation of claim ¹²²~~123~~, wherein said amount of said bioavailable source of chromium comprises no less than 200 micrograms of elemental chromium.

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~~125~~. The ingestible formulation of claim ¹²³~~124~~, wherein said amount of said bioavailable source of vanadium comprises no less than 5 mg of elemental vanadium.

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~~126~~. The ingestible formulation of claim ¹²⁴~~125~~, further comprising an effective amount of one or more of the following: aspirin, Vitamin E, and a bioavailable source of magnesium.

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127. The ingestible formulation of claim ~~124~~¹²³, wherein said anti-diabetic agent comprises a type of anti-diabetic agent selected from the group consisting of thiazolidinediones, sulfonylureas, benzoic acid derivatives, and alpha-glucosidase inhibitors.

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128. A pill for improving glucose metabolism in a subject with abnormal glucose metabolism, comprising:

(a) a bioavailable source of chromium in a complex and amount that delivers an effective amount of chromium for improving glucose metabolism; and

(b) a bioavailable source of vanadium in a complex and amount that delivers an effective amount of vanadium for improving glucose metabolism; and

(c) an anti-diabetic agent other than insulin,

wherein treatment with one or more of said pills on a daily basis for a period of at least about thirty days reduces the Hb1Ac level of said subject by at least about 10% as compared to treatment with said anti-diabetic agent alone.

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129. The pill of claim 128, wherein said amount of said bioavailable source of vanadium comprises no less than 5 mg of elemental vanadium.

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130. The pill of claim 129, wherein said amount of said bioavailable source of chromium comprises no less than 5 micrograms of elemental chromium.

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131. The pill of claim 130, further comprising an effective amount of one or more of the following: a bioavailable source of aspirin, Vitamin E, and a bioavailable source of magnesium.

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132. The pill of claim 131, wherein said anti-diabetic agent comprises a type of anti-diabetic agent selected from the group consisting of thiazolidinediones, sulfonylureas, benzoic acid derivatives, and alpha-glucosidase inhibitors.

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133. A kit for improving glucose metabolism in a subject comprising:

(a) an ingestible formulation comprising a bioavailable source of chromium in a complex and amount that delivers an effective amount of chromium for improving glucose metabolism, a bioavailable source of vanadium in a complex and amount that delivers an effective amount of vanadium for improving glucose metabolism, and an anti-diabetic agent other than insulin; and

(b) instructions for the administration of said ingestible formulation,

wherein use of one or more of said kits in accordance with said instructions by said subject for a period of at least about thirty days reduces the Hb1Ac level for said subject by at least about 10% as compared to treatment with said anti-diabetic agent alone.

133
134. The kit of claim ~~134~~¹³², wherein said instructions provide for (a) the simultaneous administration of said chromium, vanadium and anti-diabetic agent, and (b) the daily dose regiment for said kit and the duration of use one or more of said kits.